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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C07C 43/32, C08G 65/32, C11D 1/72		A1	(11) International Publication Number: WO 99/32424
			(43) International Publication Date: 1 July 1999 (01.07.99)
(21) International Application Number: PCT/SE98/02239 (22) International Filing Date: 8 December 1998 (08.12.98) (30) Priority Data: 9704755-9 19 December 1997 (19.12.97) SE (71) Applicant (for all designated States except US): AKZO NOBEL N.V. [NL/NL]; P.O. Box 9300, NL-6800 SB Arnhem (NL). (72) Inventors; and (75) Inventors/Applicants (for US only): BERGSTRÖM, Karin [SE/SE]; PI 3842, S-442 95 Kungälv (SE). HELLBERG, Per-Erik [SE/SE]; Hog 9055, S-444 97 Svenshögen (SE). (74) Agent: GOTTBERG-KLINGSKOG, Eva; Akzo Nobel Surface Chemistry AB, S-444 85 Stenungsund (SE).		(81) Designated States: AU, BR, BY, CA, CN, CZ, EE, ID, IL, IN, JP, KR, LT, LV, MX, NO, NZ, PL, RO, RU, SG, SI, UA, US, VN, Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published With international search report.	
(54) Title: AN ORTHO ESTER-BASED SURFACTANT, ITS PREPARATION AND USE			
<div style="text-align: center;"> $\begin{array}{c} \text{O} - (\text{A}_1)_{n_1}\text{R}_1 \\ \\ \text{R} - \text{C} - \text{O} - \text{R}_2 \\ \\ \text{O} - (\text{A}_2)_{n_2}\text{R}_2 \end{array} \quad (I)$ </div>			
(57) Abstract <p>The present invention relates to a new ortho ester-based surfactant, where the hydrophobic and hydrophilic parts are connected by ortho ester linkages to the molecule. The ortho ester has formula (I), where R is hydrogen or an aliphatic group with 1-7 carbon atoms; R₁ is hydrogen or an alkyl group with 1-5 carbon atoms; A₁ is an alkyleneoxy group with 2-4 carbon atoms, the number of ethyleneoxy groups being at least 50 % of the total number of alkyleneoxy groups; n₁ is a number between 1 and 30; R₂ is an aliphatic group with 5-22 carbon atoms; A₂ is an alkyleneoxy group with 3-4 carbon atoms; n₂ is a number between 0-30, provided that when R₂ is an aliphatic group with 5-6 carbon atoms n₂ is at least 1; R₃ is selected from the group consisting of (A₁)_{n1}R₁, (A₂)_{n2}R₂ and an alkyl group with 1-6 carbon atoms, where A₁, n₁, R₁, A₂, n₂ and R₂ have the same meaning as mentioned above, or a di- or polycondensate via any of the free hydroxy groups of the ortho ester. The ortho ester surfactants are stable in alkaline solutions, but are readily hydrolysed in acidic solutions to yield products that are not surface active. They are suitable to be used as emulsifiers or dispersants.</p>			

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AN ORTHO ESTER-BASED SURFACTANT, ITS PREPARATION AND USE.

The present invention relates to a new ortho ester-based surfactant, where the hydrophobic and hydrophilic parts are connected by ortho ester linkages to the molecule. The surfactants are stable in alkaline solutions, but are readily hydrolysed in acidic solutions to yield products that are not surface active.

Surfactants are used in a variety of applications and processes, but once their task is fulfilled their presence is often undesirable. From an environmental point of view, it is a great advantage if products that ultimately end up in the environment are easily degradable, either biologically or by other means. Also, since surfactants have the ability to form emulsions and dispersions, which in most cases is the very reason for using them, they make the separation of hydrophobic material from the waste water obtained in industrial processes difficult.

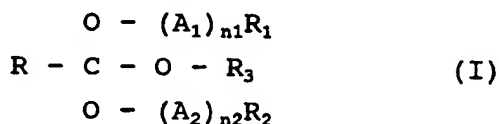
In order to improve the degradability of surfactants and make the separation of hydrophobic material from waste water easier, it has been suggested in EP-A1-0 742 177 and EP-A1-0 742 178 to use hydrolysable aldehyde- and ketone-based surfactants. The surfactants, which contain acetal linkages, are stable in alkaline solutions but are hydrolysed in acidic solutions. Acetal-based surfactants are also described in EP-A3-0 054 366.

However, to effect complete hydrolysis the pH need to be lower and the reaction time longer for the acetals as compared to the ortho esters. This will result in a larger consumption of chemicals, and either give a water phase with an unacceptably low pH to be let out to the sewage treatment works or, if the waste water is neutralised, the formation of larger amounts of salt. Furthermore, there is only a small number of long-chain aldehydes that are commercially available, and consequently the range of acetal-based surfactants possible to obtain is limited. In addition the aldehydes are generally more difficult to produce than the

corresponding alcohols, and are therefore more expensive.

Ortho ester surfactants have been described in EP-A1-564 402, where an ortho ester group is used for end-capping of nonionic surfactants. The products obtained are low-foaming, and can be used e.g. in machine dish-washing and bottle-cleaning. These products, however, will only marginally benefit from a better degradation, since a hydrolysis step will produce compounds that are still surface active.

The aim of the present invention is to provide surfactants with at least as good emulsifying and dispersing ability as conventional types of surfactants, which in addition are readily cleavable and more easily biodegradable. Their degradation products should also be environmentally friendly, and not exhibit any essential surface activity. Further, this new kind of surfactants should be easy to produce. Surprisingly, it has now been found that surfactants based on an ortho ester according to formula



where R is hydrogen or an aliphatic group with 1-7 carbon atoms; R₁ is hydrogen or an alkyl group with 1-5 carbon atoms, preferably R₁ is an alkyl group with 1-4 carbon atoms; A₁ is an alkyleneoxy group with 2-4 carbon atoms, the number of ethyleneoxy groups being at least 50% of the total number of alkyleneoxy groups; n₁ is a number between 1 and 30, preferably between 2-25; R₂ is an aliphatic group with 5-22 carbon atoms, preferably 8-22; A₂ is an alkyleneoxy group with 3-4 carbon atoms; n₂ is a number between 0-30, preferably between 0-20, provided that when R₂ is an aliphatic group with 5-6 carbon atoms n₂ is at least 1, preferably at least 2; R₃ is selected from the group consisting of (A₁)_{n₁}R₁, (A₂)_{n₂}R₂ and an alkyl group with 1-6 carbon atoms, preferably 1-4, where A₁, n₁, R₁, A₂, n₂ and R₂ have the same meaning as mentioned above; or a di- or polycondensate via any of the free hydroxy groups of the ortho-ester; display the above-mentioned properties. To

strengthen the hydrophilic part of the molecule the groups A_1 may consist only of ethyleneoxy groups.

5 The surfactants of formula I have a good emulsifying and dispersing ability, and are preferably used in applications where the rapid cleavability offers an advantage, e.g. for hard surface cleaning, deinking, viscose processing, disinfection and in fibre and textile processes such as dyeing and scouring. They are also low-foaming, which is an advantage in many applications. When used for cleaning of
10 hard surfaces they exhibit comparable or better effects than traditional surface active nonionic alkylene oxide adducts. The emulsifying ability of this surfactant type is further demonstrated by the formulation of an emulsion for a pesticide. This formulation is of comparable stability to a
15 formulation obtained with an optimized nonionic emulsifier of the traditional type.

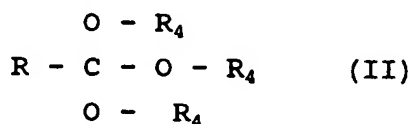
The cleavage of the ortho ester-based surfactant is promoted to a high degree by decreasing the pH and increasing the temperature. The ortho ester could also be used as
20 an emulsifier/dispersant at a lower pH, e.g. at a pH of 5, if the process is fast enough, and consequently be cleaved at the same pH. When compared to the cleavage of an acetal-based surfactant at the same conditions, the ortho ester-based surfactant is cleaved much more rapidly. The cleavage
25 results in degradation products that lack the ability to behave as surfactants, e.g. to form emulsions, which is demonstrated in Example 12. The rapid cleavability of the ortho esters of the present invention presents a special advantage in application areas where the separation of an
30 oil phase from the water phase is desirable, e.g. for waste water treatment, in the working-up of emulsions formed when cleaning hard surfaces, and in deinking and textile processes.

35 The invention also relates to a process for making the ortho ester-based surfactants, where low molecular weight ortho esters are used as starting materials. These
~~low molecular weight ortho esters are reacted with a hydro-~~

phobic component, which is an alcohol, and preferably an end-capped hydrophilic component, which is preferably a polyethylene oxide adduct. The molar amounts of the reactants are preferably 1-2 moles of the hydrophilic component per mole orthoester and 1-2 moles of the hydrophobic component per mole ortho ester.

By this process surface active ortho esters are obtained, where the hydrophobic and hydrophilic parts each individually is connected by ortho ester bonds to the molecule.

The ortho ester-based surfactants of the present invention can be produced by reacting an ortho ester of the general formula



where R has the same meaning as in Formula I and R_4 is an alkyl group with 1-6 carbon atoms, preferably 1-4, in one or several steps, with reactants having the formulas $\text{HO}(\text{A}_1)_{n_1}\text{R}_1$ and $\text{HO}(\text{A}_2)_{n_2}\text{R}_2$, where R_1 , R_2 , A_1 , A_2 , n_1 and n_2 have the same meaning as in Formula I, while evaporating alcohols with the formula R_4OH , where R_4 has the same meaning as above. The reaction preferably is performed in the presence of an acid, e.g. methanesulphonic acid, p-toluenesulphonic acid or citric acid. The temperature is increased during the reaction and is finally reaching 140 to 220°C. The alcohols R_4OH , that are liberated during the reaction, are gradually evaporated from the reaction mixture. In the final phase of the reaction vacuum is applied to remove the residual amounts of alcohols, thereby driving the reaction to completion.

Suitable examples of ortho esters II are methyl or ethyl ortho formate, methyl or ethyl ortho acetate and other low molecular weight ortho esters that are commercially available.

The hydrophobic part of the molecule may be derived from an alcohol R_2OH , or an alkoxylate thereof. The alcohol

could either be synthetic or natural. Suitable examples of alkyl groups R_2 are 2-ethylhexyl, octyl, decyl, coco alkyl, lauryl, oleyl, rape seed alkyl and tallow alkyl. Other suitable hydrocarbon groups R_2 are those obtained from oxoalcohols, Guerbet alcohols, and methyl substituted alcohols with 2-4 groups having the formula $-\text{CH}(\text{CH}_3)-$ included in the alkyl chain. The alcohols may also be propoxylated or butoxylated.

The hydrophilic part of the molecule is preferably derived from polyethylene glycols, that are end-capped, preferably with a methyl or ethyl group, and have a molecular weight between 100 and 2000. The choice of hydrophobic and hydrophilic parts and the relative amounts of them will of course vary between different applications, to satisfy their demands for a specific HLB, cloud point etc.

In another embodiment the present invention relates to the use of an ortho ester according to formula I in a process comprising

- a) emulsifying or dispersing a hydrophobic component in water at a pH of 6 or above, preferably at pH 7 or above, in the presence of an ortho ester according to the present invention,
- b) lowering the pH or increasing the temperature of the emulsion or dispersion, or a combination thereof, and thereby breaking the emulsion or dispersion, and
- c) separating the hydrophobic component from water.

The surfactant is normally used as an emulsifier or dispersant at a pH of 9 or above, but could also be used down to a pH of ca 6. The waste water obtained as a result of the performance of the surfactant in step a is then treated in accordance with b. The pH of the emulsion or dispersion is preferably reduced to a pH of between 4 and 6. If needed, the temperature may be raised, preferably to between 20 and 60°C to further promote the cleavage. In some circumstances, when the pH during the emulsification is low enough, it might suffice to raise the temperature. The lower the pH and the higher the temperature, the faster the

cleavage will occur. In most circumstances it might be more convenient to lower the pH further than to raise the temperature above ambient, since the latter will often require a large energy input.

5 The above-mentioned process could be used in a variety of applications. One major application is the cleaning of hard surfaces, for example in connection with vehicle cleaning and the cleaning of storage tanks and tankers, where the ortho ester surfactant is used at alkaline pH as
10 an emulsifier or dispersant for the hydrophobic dirt or fluid. When the surface has been cleaned, the waste water is acidified whereby the surfactant is cleaved. This causes the emulsion or dispersion to break, and the hydrophobic material is separated from the water phase.

15 In an analogous manner, the hydrophobic ink obtained in a deinking process, the surplus of hydrophobic dye from a textile dyeing process and the dirt from a textile scouring process could be emulsified or dispersed by the surfactants, and later on removed from the process waste water.

20 The ortho ester surfactants also benefit from a better biodegradability than the corresponding conventional nonionic surfactants. When subjected to a neutral or slightly acidic pH in a sewage-treatment plant, the ortho ester surfactants are cleaved to yield non-toxic substances
25 that are essentially not surface active. These substances would be more easily biodegraded than an intact surface active molecule would be. A comparison between a traditional nonionic surfactant and an ortho ester surfactant according to the present invention (see Example 14) shows the latter
30 to be more easily biodegradable.

The present invention is further illustrated by the following Examples.

Example 1

35 1.5 moles of a C₉-C₁₁ linear primary alcohol (Dobanol 91 from Shell), 1.5 moles of diethylene glycol monoethyl ether ethoxylate (diethylene glycol monoethyl ether + 2 moles of ethylene oxide), 1 mole of triethyl orthoformate

and 0.2% w/w of methanesulphonic acid were mixed together at ambient temperature. The temperature of the reaction mixture was gradually raised, and finally, after ca 4 hours, a temperature of 150-200°C was reached. Ethanol, that was liberated during the reaction, was continuously distilled off. In the final phase of the reaction, vacuum was applied to facilitate the removal of ethanol. A total of 30.8 g of ethanol was collected, which corresponds to 92% of the theoretical amount. The distillate was analysed by ¹H-NMR and the product was analysed by ¹H-NMR and ¹³C-NMR. According to the analyses there was no unreacted triethyl orthoformate left.

Example 2

1.05 moles of 2-ethylhexanol propoxylate (2-ethylhexanol + 13 moles of propylene oxide), 1.05 moles of diethylene glycol monomethyl ether ethoxylate (diethylene glycol monomethyl ether + 18 moles of ethylene oxide), 1.0 mole of triethyl orthoformate and 1% w/w of anhydrous citric acid were mixed and a reaction was performed under the same conditions as in Example 1. A total of 10.2g of ethanol was distilled off, which corresponds to 92% of the theoretical amount. The same analyses as in Example 1 were performed. No unreacted triethyl orthoformate was found.

Example 3

1.5 moles of 2-ethylhexanol, 1.5 moles of a monomethyl-blocked polyethylene glycol having a mean molecular weight of 550, 1 mole of triethyl orthoformate and 0.2% w/w of methanesulphonic acid were mixed. The same procedure as in Example 1 was followed. A total of 18.2 g of ethanol was collected, which corresponds to 98% of the theoretical amount. The same analyses as in Example 1 were performed. No unreacted triethyl orthoformate was found. According to the NMR analysis more than 70% of the product consists of three surface active components specified below. The number of ethoxy groups that has not been substituted is denoted by x, the number of 2-ethylhexyl groups by y and the number of end-capped polyoxyethylene-groups by z.

Component nr	x	y	z
1	1	1	1
2	0	2	1
3	0	1	2

Example 4

The procedure is the same as in Example 1, except that 2-ethylhexanol + 2.4 moles of propylene oxide and monomethyl-blocked polyethylene glycol having a mean molecular weight of 550 are used as the hydrophobic and hydrophilic components respectively. The same analyses as in Example 1 were performed. No unreacted triethyl orthoformate was found.

Example 5

The procedure is the same as in Example 1, except that n-octanol and monomethyl-blocked polyethylene glycol having a mean molecular weight of 350 are used as the hydrophobic and hydrophilic components respectively. The same analyses as in Example 1 were performed. No unreacted triethyl orthoformate was found.

Example 6

0.152 moles of hexadecanol, 0.152 moles of monomethyl-blocked polyethylene glycol having a mean molecular weight of 750, 0.101 moles of triethyl orthoformate and 0.15% w/w of anhydrous citric acid were mixed together. The temperature of the reaction mixture was gradually raised from 22°C to 155°C during the course of 30 min. Ethanol, that was liberated during the reaction, was continuously distilled off. When the temperature had reached 155°C, the pressure was slowly reduced to 3 mbar and held there for 20 min to facilitate the removal of ethanol. A total of 13.7 g of ethanol was collected, which corresponds to 99% of the theoretical amount. The distillate was analysed by ¹H-NMR and the product was analysed by ¹H-NMR and ¹³C-NMR. According to the analyses there was no unreacted triethyl orthoformate left, and 60% of the product mixture consists of three sur-

face active components corresponding to the ones specified in the table of Example 3.

Example 7

5 The procedure is the same as in Example 6, except that 2-ethylhexanol + 2 moles of propylene oxide and monomethyl-blocked polyethylene glycol having a mean molecular weight of 550 are used as the hydrophobic and hydrophilic components respectively. The same analyses as in Example 6 were performed. No unreacted triethyl orthoformate was found.

Example 8

15 The procedure is the same as in Example 6, except that a C₉-C₁₁ linear primary alcohol (Dobanol 91 from Shell) and monomethyl-blocked polyethylene glycol having a mean molecular weight of 470 are used as the hydrophobic and hydrophilic components respectively. The same analyses as in Example 6 were performed. No unreacted triethyl orthoformate was found.

Example 9

20 The biodegradability of the ortho ester-based surfactant described in Example 1 was investigated by the "closed bottle test" as described in OECD Test 301D. The surfactant reached 82% biodegradation after 28 days, and is consequently classified as easily biodegradable.

Example 10

25 To evaluate the cleaning efficiency of formulations containing an ortho ester-based surfactant of the present invention, the following cleaning test was used: Alumina plates (41x15mm) coated with C14 marked triolein (glycerol trioleate from Amersham) were put in a holder and washed in a Terg-O-Tometer by turning the holders back and forth through the surfactant solution with a velocity of 50 rpm during a period of 5 min. The test was performed at 20 and 40°C, using the formulations I, II and III in the Table below, where I is a reference formulation containing a traditional nonionic surfactant. The formulations were diluted with tap water 1:100 prior to use. After the cleaning step

the holders were dipped for 5 seconds in tap water holding a temperature of 20°C.

Component	Formulation I % by weight of component	Formulation II % by weight of component	Formulation III % by weight of component
C ₉ -C ₁₁ alcohol + 4EO	5	-	-
Ortho ester mixture of Ex. 4	-	5	-
Ortho ester mixture of Ex. 7	-	-	5
Octyliminodi- propionate	2.8	2.8	2.8
Tetrapotassium pyrophosphate	6	6	6
Metasilicate	4	4	4
Water	balance	balance	balance

The plates were transferred to vials containing scintillation liquid (Ultima Gold from Packard) with the ability to dissolve fat, and were shaken at 300 rpm for 20 minutes. The plates were removed from the vials, and the liquid was analysed for radioactivity in a scintillation analyser (Tri-Carb 1900CA from Packard). The result is reported as % washed-away fat as compared to coated plates that have not been washed according to the procedure above.

Formulation	% washed-away fat at 20°C	% washed-away fat at 40°C
I	57.9	85.0
II	87.6	79.6
III	84.2	91.2

This example shows that the ortho ester surfactants

are as good cleaners for hard surfaces as the nonionic surfactant used as a reference.

Example 11

The wetting ability of ortho ester-based surfactants according to the present invention was estimated by contact angle measurements.

The test solutions were the formulations I (reference), II and III that were used in Example 10. The formulations were used undiluted. The contact angle was measured against a hydrophobic polymeric material (Parafilm PM-992 from American Can Company) with a Ramé-Hart NRL C.A. Goniometer. The measurements were made after 1 min, and each value is the average of 10 measurements. The results are summarised in the Table below.

Formulation	Contact angle after 1 min (°)
I	35.8
II	35.3
III	34.4

The results show that the wetting ability of the ortho ester-based surfactants according to this invention is comparable to the wetting ability of the reference compound.

Example 12

A comparison is made between the ortho ester-based surfactant produced according to Example 1, an ortho ester end-blocked surfactant and an acetal-based surfactant, regarding their emulsifying ability at different pH-values.

The oil phase used in all experiments is n-decane, and the water phase consists of different buffer solutions. Sudan Red B is added as a colorant for the emulsion. The surfactants used are:

- A) Ortho ester-based surfactant according to Example 1
- B) A C₁₀-C₁₁ linear primary alcohol + 8 moles of ethylene oxide that is end-blocked with triethyl orthoformate (synthesized according to the procedure described in

EP-A1-0 564 402)

- C) The acetal between n-decanal and glycerol ethoxylated with 4 moles of ethylene oxide (synthesized according to the procedure described in EP-A1-0 742 177)

Procedure:

7.5 ml n-decane, 7.5 ml buffer solution, 0.3 g surfactant and 2 drops of Sudan red B are placed in a sealable test tube, which is manually shaken for one minute before the first measurement. The emulsion is allowed to separate, and after 3 minutes the degree of separation is measured as $a/b \times 100$, where a is mm clear lower phase, and b is mm clear water phase before emulsifying.

Between the measurements the test tubes are continuously shaken at 1000 rpm in the horizontal position by an IKA-VIBRAX-VXR apparatus. Before each new measurement the test tube is manually shaken for 30 s to completely reemulsify the mixture.

The test is performed at 22 and 50°C. For the test at 50°C n-decane and buffer solution is preheated before the surfactant is added.

pH	Time for 100% separation of emulsions with A at 22°C	Time for 100% separation of emulsions with A at 50°C	Time for 100% separation of emulsions with C at 22°C
2	< 5 minutes	< 1 minute	> 35 days
3	< 5 minutes	< 1 minute	-
4	< 39 minutes	< 10 minutes	-
5	< 120 minutes	< 15 minutes	-
6	ca 55 hours	< 3 hours	-
7	> 8 days	< 22 hours	-
8	> 15 days	ca 29 hours	-
9	-	> 16 days	-
10	> 40 days	> 40 days	-
11.5	> 40 days	> 40 days	-

- test not performed

At 22°C surfactant A is readily hydrolysed at pH-values below 5, which results in a loss of surface activity causing the emulsions to separate. At 50°C the hydrolysis is rapid in this pH range, yielding separation times below 15 minutes. As expected, the hydrolysis is slow at alkaline pH.

To obtain an acceptable hydrolysis time for C, 20% H₂SO₄ was added. This resulted in more than 3 hours time for 100% separation, whereas for an addition of 5% H₂SO₄, more than 72 hours was needed.

The emulsion produced with surfactant B is only marginally effected by pH. Still after 11 days at a pH of 2 and 22°C, only a separation of 11% is obtained. This is expected since the surface activity of B does not disappear when the ortho ester bond is broken.

These above results show the superiority of surfactant A to surfactant B and C regarding ease of hydrolysis at acid pH-values.

Example 13

To further investigate the emulsion separation obtained at a pH of 5, two other kinds of oils were emulsified by the same surfactants A and C that were used in Example 8. For a comparison with a conventional nonionic surfactant, a C₉-C₁₁ alcohol + 4 moles of ethylene oxide was used (surfactant D). The oils used were refined soy bean oil (produced by Karlshamn) and diesel oil, and the water phase consists of a pH 5 buffer solution .

Procedure:

At room temperature 200 ml oil, 300 ml buffer solution and 6 g surfactant are placed in a 500 ml reactor equipped with a mechanical stirrer of the propeller type. The reactor has an outlet at the bottom. The mixture is vigorously stirred at ca 500 rpm for 90 minutes. The emulsion that is formed is allowed to stand for 5 minutes, after which 250 ml liquid is drained off through the bottom outlet during ca 1 minute. The sample is left for 3 hours, after which the volume of oil and/or emulsion is measured. The result is presented as % v/v referring to the total volume

of the sample. The values obtained are collected in the table below.

Surfactant	pH	oil	% oil/emulsion in sample
A	5	soy bean	4.0
C	5	soy bean	35.5
D	5	soy bean	51.6
A	5	diesel	0.8
C	5	diesel	75.9
D	5	diesel	72.8

The amount of oil and/or emulsion in a sample reflects the rate of hydrolysis of the surfactants. The results obtained show clearly that the use of the cleavable ortho ester-based surfactant A facilitates the separation of the oil phase from emulsions containing vegetable oil as well as petroleum based oil. The rate of emulsion separation at pH 5 is substantially higher using this ortho ester-based surfactant as compared to using the acetal-based surfactant C. When using the conventional nonionic surfactant D the rate of separation is still slower.

Example 14

A formulation for a pesticide containing an ortho ester-based surfactant is compared to a standard formulation containing a traditional nonionic surfactant.

Component	Formulation I amount of component (g/l)	Formulation II amount of component (g/l)
Dimethoate	200	200
Ortho ester mixture of Example 2	50	-
C ₈ -alcohol alkoxylate (EO/PO), MW 1800	-	50
Cyclohexanone	301	301
Xylene	413	413

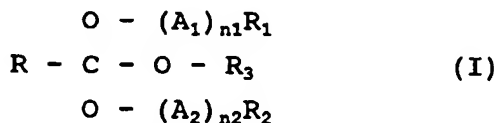
5 ml of each formulation was emulsified in 95 ml of water, and the emulsions were transferred to 100 ml test tubes. The separation of the emulsions were noted as % v/v of clear upper phase at certain time intervals. The results are summarized in the table below.

Formulation	Time	Separation (% v/v)
I	30 minutes	0
I	1 hour	0.5
I	2 hours	0.5
II	30 minutes	0
II	1 hour	0
II	2 hours	0

The formulation containing the ortho ester-based surfactant yielded an emulsion that was of comparable stability to the emulsion obtained by formulation II, which contains a traditional nonionic surfactant. However, the ortho ester-based surfactant has the advantage to be more easily cleaved to non-surface active compounds, and has therefore a better environmental profile than the conventional nonionic surfactant. The biodegradability of the ortho ester-based surfactant according to the "closed bottle test" was 37% after 28 days and 41% after 42 days, whereas for the corresponding conventional surfactant 18% was degraded after 28 days and 35% after 112 days.

C L A I M S

1. An ortho ester **characterized in**, that it has the formula



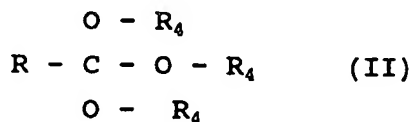
where R is hydrogen or an aliphatic group with 1-7 carbon atoms; R₁ is hydrogen or an alkyl group with 1-5 carbon atoms; A₁ is an alkyleneoxy group with 2-4 carbon atoms, the number of ethyleneoxy groups being at least 50% of the total number of alkyleneoxy groups; n₁ is a number between 1 and 30; R₂ is an aliphatic group with 5-22 carbon atoms; A₂ is an alkyleneoxy group with 3-4 carbon atoms; n₂ is a number between 0-30, provided that when R₂ is an aliphatic group with 5-6 carbon atoms n₂ is at least 1; R₃ is selected from the group consisting of (A₁)_{n₁}R₁, (A₂)_{n₂}R₂ and an alkyl group with 1-6 carbon atoms, where A₁, n₁, R₁, A₂, n₂ and R₂ have the same meaning as mentioned above; or a di- or poly-condensate via any of the free hydroxy groups of the ortho ester.

2. An ortho ester according to claim 1, where R₁ is an alkyl group with 1-4 carbon atoms.

3. An ortho ester according to claims 1-2, where n₁ is a number between 2-25 and n₂ is a number between 0-20.

4. An ortho ester according to claims 1-3, where n₂ is 0, R₂ is an aliphatic group with 8-22 carbon atoms and A₁ is an ethyleneoxy group.

5. A process for the preparation of an ortho ester according to any of the claims 1-4, **characterized in**, that it is prepared by reacting an ortho ester of the general formula



where R has the same meaning as in any of the claims 1-4 and R₄ is an alkyl group with 1-6 carbon atoms, in one or

several steps, with reactants having the formulas $\text{HO}(\text{A}_1)_{n_1}\text{R}_1$ and $\text{HO}(\text{A}_2)_{n_2}\text{R}_2$, where R_1 , R_2 , A_1 , A_2 , n_1 and n_2 have the same meaning as in any of the claims 1-4, while evaporating alcohols with the formula R_4OH , where R_4 has the same meaning as above.

6. Use of an ortho ester having the formula I, as defined in claims 1-4, as an emulsifying and/or dispersing agent.

7. Use of the ortho ester according to claim 6 in a cleaning, scouring, dyeing and deinking process and in the formulation of a pesticide composition.

8. Use of an ortho ester according to claims 6 or 7 in a process comprising

- a) emulsifying or dispersing a hydrophobic component in water at a pH of 6 or above in the presence of an ortho ester in accordance with claims 1-4,
- b) lowering the pH or increasing the temperature of the emulsion or dispersion, or a combination thereof, and thereby breaking the emulsion or dispersion, and
- c) separating the hydrophobic component from water.

9. Use of an ortho ester in a process according to claim 8, where the temperature in step b is raised to between 20 and 60°C.

10. Use of an ortho ester in a process according to claims 8-9, where the pH in step b is between 4 and 6.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 98/02239

A. CLASSIFICATION OF SUBJECT MATTER		
IPC6: C07C 43/32, C08G 65/32, C11D 1/72 According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols)		
IPC6: C07C, C08G, C11D		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
SE,DK,FI,NO classes as above		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
E	WO 9800452 A1 (AKZO NOBEL N.V.), 8 January 1998 (08.01.98) --	1-10
A	EP 0564402 A1 (CIBA-GEIGY AG), 6 October 1993 (06.10.93) --	1-10
A	EP 0054366 A2 (IMPERIAL CHEMICAL INDUSTRIES PLC), 23 June 1982 (23.06.82) --	1-10
A	US 4450087 A (HERBERT F. ASKEW ET AL), 22 May 1984 (22.05.84), See part. paragraph d in the description, column 2 --	1-10
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input checked="" type="checkbox"/> See patent family annex.		
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family		
Date of the actual completion of the international search		Date of mailing of the international search report
19 March 1999		22-03- 1999
Name and mailing address of the ISA:		Authorized officer
Swedish Patent Office Box 5055, S-102 42 STOCKHOLM Facsimile No. + 46 8 666 02 86		Eva Johansson Telephone No. + 46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 98/02239

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 3903006 A (JOHN SCOTCHFORD ELLIOTT ET AL), 2 Sept 1975 (02.09.75) --	1-10
A	US 2867667 A (OTIS C. DERMER ET AL), 6 January 1959 (06.01.59) -- -----	1-10

INTERNATIONAL SEARCH REPORT

Information on patent family members

02/03/99

International application No.

PCT/SE 98/02239

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